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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/785,220	02/24/2004	Avi Ashkenazi	P1216R1C1D4	1253
9157	7590 02/25/2005		EXAMINER	
GENENTECH, INC.			HADDAD, MAHER M	
I DNA WAY SOUTH SAN	FRANCISCO, CA 94080)	ART UNIT	PAPER NUMBER
,			1644	

DATE MAILED: 02/25/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

	_	Application No.	Applicant(s)			
Office Action Summary		10/785,220	ASHKENAZI ET AL.			
		Examiner	Art Unit			
		Maher M. Haddad	1644			
Period fo	The MAILING DATE of this communication app or Reply	ears on the cover sheet with the c	orrespondence address			
THE - External after - If the - If NC - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY MAILING DATE OF THIS COMMUNICATION. nsions of time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. period for reply specified above is less than thirty (30) days, a reply period for reply is specified above, the maximum statutory period were to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing ed patent term adjustment. See 37 CFR 1.704(b).	86(a). In no event, however, may a reply be time within the statutory minimum of thirty (30) days ill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).			
Status						
2a)	Responsive to communication(s) filed on This action is FINAL. 2b) This Since this application is in condition for allowant	action is non-final.	secution as to the merits is			
	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.					
Dispositi	on of Claims	•				
4) ☐ Claim(s) 49-63 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration. 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) 49-63 are subject to restriction and/or election requirement.						
Applicati	on Papers	·				
9)[The specification is objected to by the Examiner					
10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.						
	Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).					
11)	Replacement drawing sheet(s) including the correction. The oath or declaration is objected to by the Exa	• • • • • • • • • • • • • • • • • • • •				
Priority u	ınder 35 U.S.C. § 119					
12)[a)[Acknowledgment is made of a claim for foreign All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priori application from the International Bureau see the attached detailed Office action for a list of	have been received. have been received in Application ty documents have been receive (PCT Rule 17.2(a)).	on No d in this National Stage			
Attachment	• •	A) 🔲 Intention Commence	(DTO 442)			
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 4) Interview Summary (PTO-413) Paper No(s)/Mail Date						
	nation Disclosure Statement(s) (PTO-1449 or PTO/SB/08) No(s)/Mail Date	5)	atent Application (PTO-152) <u>tion Sheet</u> .			

Continuation of Attachment(s) 6). Other: Notice to comply with sequence requirements.

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DETAILED ACTION

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Sequence Compliance

- 1. There is a discrepancy in the length of SEQ ID NO: 11. Figure 5 shows that SEQ ID NO: 11 is a 1842 nucleotide sequence. The sequence listing indicates that SEQ ID NO:11 is a 2181 nucleotide sequence. Clarification is required.
- 2. The specification on page 49, lines 36-40, discloses that clone DNA40628 contains a single open reading frame with an apparent translational initiation site at nucleotide positions 52-54 (FIG. 5; SEQ ID NO: 11). The sequence listing SEQ ID NO:11 does not correspond with the translation initiation site at nucleotide positions 52-54. Clarification is required.
- 3. This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 CFR 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 CFR 1.821 through 1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures.

The specification contains the nucleic acid sequence of 1,842 bp on Fig.5, which is not present in sequence listing. While the sequence identifier indicates that said nucleic acid is SEQ ID NO:11, the sequence listing indicates that SEQ ID NO: 11 is 2181 nucleotide sequence.

Restriction/Election

- 4. Given the apparent ambiguity about SEQ ID NO:11 and its relationship with SEQ ID NO:1, the cDNA deposited under ATCC accession number 209432 and PRO301 polypeptide, the claims have been limited to either an isolated nucleic acid molecule encoding the polypeptide of SEQ ID NO:1, or an isolated nucleic acid sequence of SEQ ID NO: 11, irrespective of the format of the claims. Due to the ambiguity, the cDNA deposited under ATCC accession number 209432 has been placed in both Groups.
- 5. Restriction to one of the following inventions is required under 35 U.S.C. § 121:
 - I. Claims 49-63, drawn to an isolated nucleic acid sequence encoding the polypeptide of SEQ ID NO:1 with or without signal sequence, and the full-length coding sequence of the cDNA deposited under ATCC accession number 209432; vectors, host cells, and methods of producing the polypeptide, classified in Class 536, subclass 23.5; Class 435, subclasses 69.1, 455, 252.3, and 320.1.

Art Unit: 1644

- II. Claims 49-63, drawn to an isolated nucleic acid sequence of SEQ ID NO:11, the full-length coding sequence of the nucleic acid sequence of SEQ ID NO: 11 and the full-length coding sequence of the cDNA deposited under ATCC accession number 209432; vectors, host cells, and methods of producing the polypeptide, classified in Class 536, subclass 23.5; Class 435, subclasses 69.1, 455, 252.3, and 320.1.
- 6. Groups I and II. Two different nucleic acids differ with respect to their structures (apparently en c coding for two different proteins) and physicochemical properties; therefore each product is patentably distinct.
- 7. These inventions are distinct for the reasons given above. Further, even though the classification is shared, a different field of search would be required based upon the structurally distinct products recited and the various methods of use comprising distinct method steps. Therefore restriction for examination purposes as indicated is proper. Further, a prior art search also requires a literature search. It is an undue burden for the examiner to search more than one invention.
- 8. Applicant is advised that the response to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed.
- 9. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Maher Haddad whose telephone number is (571) 272-0845. The examiner can normally be reached Monday through Friday from 7:30 am to 4:00 pm. A message may be left on the examiner's voice mail service. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Chan can be reached on (571) 272-0841. The fax number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Maher Haddad, Ph.D.

Maker Haddad

Patent Examiner

Technology Center 1600

Application No.: 10/785, 220

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

X	 This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
	 This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
	A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
	4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
	5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
	6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
囟	7. Other: Fig 5 Contains a nucleic acid that is not present in sequence listing
Ap	plicant Must Provide:
X	An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
囚	An initial or <u>substitute</u> paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
Ø	A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).
Fo	r questions regarding compliance to these requirements, please contact:
Fo	or Rules Interpretation, call (703) 308-4216 or CRF Submission Help, call (703) 308-4212
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For Patentin software help, call (703) 308-6856